The Journal of Molecular Diagnostics, Vol. 16, No. 3, May 2014

the Journal of Molecular Diagnostics

jmd.amjpathol.org

CORRECTION

In the article entitled, "A Clinically Validated Diagnostic Second-Generation Sequencing Assay for Detection of Hereditary *BRCA1* and *BRCA2* Mutations" (Volume 15, pages 796–809 of the November 2013 issue of *The Journal of Molecular Diagnostics*), Supplemental Tables S1 and S2 contained errors. Incorrect Human Genome Variation Society (HGVS) nomenclature was used to describe 10 variants. To correct those errors, 9 variants (of 398) were updated in Supplemental Table S1 (retrospective samples), and 1 variant (of 517) was updated in Supplemental Table S2 (prospective samples). In addition, the protein HGVS nomenclature was updated to use 3-letter amino acid abbreviations. The conclusions of the paper (eg, the ability to detect variants and concordance) are not altered by the incorrect notation. The corrected Supplemental Tables S1 and S2 appear online. The authors apologize for the error.

Supplemental Data

Supplemental material for this article can be found at *http://dx.doi.org/10.1016/j.jmoldx.2014.03.001*.

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